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Substitute for form 1449A/PTO		Complete if Known	
INFORMATION DISCLOSURE STATEMENT BY APPLICANT <i>(Use as many sheets as necessary)</i>		Application Number	10/573,890
		Filing Date	03-29-08
		First Named Inventor	Kazutaka Nakamoto
		Art Unit	1625
		Examiner Name	Patricia L. Morris
Sheet	1	of	3
Attorney Docket Number			

U.S. PATENT DOCUMENTS

FOREIGN PATENT DOCUMENTS

Examiner Initial *	Cite No. 1	Foreign Patent Document		Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, columns, Lines, Where Relevant Passages or Relevant Figures Appear	Text
		Country ³	Number ⁴ Kind Code (if known) ⁵				
	12	EP	0 124 154	11-07-1984			
	13	WO	03-091226	11-06-2003			
	14	WO	03-091227	11-06-2003			
	15	JP	2001-527083	12-25-2001			X
	16	WO	2004-014366	02-19-2004			
	17	JP	2005-528751	09-08-2005			X
	18	JP	2006-519247	08-24-2006			X
	19	WO	2009-084621	07-09-2009			X
	20	JP	5-284935	11-09-1993			X
	21	JP	59-073579	04-25-1984			X

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3. Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3.1.4). For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. 5. Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST 6. 16 is possible. 6. Amendment is in place a check mark here if English language Translation is attached

To the best of my knowledge, this application is in English language. Translation is attached.
This collection of information is required by 37 CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 2 hours to complete, including gathering, preparing, and submitting the completed application form the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, P.O. Box 1450 Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS.

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Substitute for form 1449A/PTO				Complete if Known	
INFORMATION DISCLOSURE STATEMENT BY APPLICANT				Application Number	10/573,890
(Use as many sheets as necessary)				Filing Date	03-29-06
				First Named Inventor	Kazutaka Nakamoto
				Art Unit	1625
				Examiner Name	Patricia L. Morris
Sheet	2	of	3	Attorney Docket Number	3939-0118PUS1

U.S. PATENT DOCUMENTS

FOREIGN PATENT DOCUMENTS

Examiner Initials ¹	Cite No. 1	Foreign Patent Document		Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Country ²	Number ³ and Code (if known) ³			
	22	JP	8-175993	07-09-1996		X

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3. Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3) 4. For Japanese patent documents, the indication of the year of the reign of the
Emperor must precede the serial number of the patent document. 5. Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST.5
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Substitute for form 1449B/PTO

INFORMATION DISCLOSURE
STATEMENT BY APPLICANT

(Use as many sheets as necessary)

Sheet

3

of

3

Complete if Known

Application Number	10/573,890
Filing Date	03-29-06
First Named Inventor	Kazutaka Nakamoto
Group Art Unit	
Examiner Name	

Attorney Docket Number 3939-0118PUS1

NON PATENT LITERATURE DOCUMENTS

Examiner initial *	Cite No. 1	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T <input checked="" type="checkbox"/>
	23	CHANG et al., "Synthesis and Structure-Activity Relationships of Quaternary Ammonium Cephalosporins with 3-Pyrazolylpyridinium Derivatives," <i>Bioorganic & Medicinal Chemistry Letters</i> (2000) Vol. 10, No. 11, pp. 1211-1214	
	24	CONNORS et al., "Prodrugs in medicine," <i>Overview, Biologicals & Immunologicals, Exp. Opin. Ther. Patents</i> , Vol. 5, No. 9, 1995, pp. 873-885	
	25	Copy of an Office Action from co-pending U.S. Patent Application No. 11/589,126, mailed May 7, 2009	
	26	HATA, "New Approaches to Antifungal Drugs for the Treatment of Fungal and Protozoal Infections, RAVUCONAZOLE and Beyond: New Targets and Pre-clinical Strategies," The SMI's 12th Annual Conference, Superbugs and Superdrugs, March 18, 2010, Crowne Plaza London - St. James, 44 pages	
	27	International Search Report dated May 20, 2008 for corresponding International Application No. PCT/JP2008/057351	
	28	ISHIKAWA et al., "TAK-599, a Novel N-Phosphono Type Prodrug of Anti-MRSA Cephalosporin T-91825: Synthesis, Physicochemical and Pharmacological Properties," <i>Bioorganic & Medicinal Chemistry</i> , Vol. 11, pp. 2427-2437, (2003)	
	29	LUKEVICS et al., "Synthesis and cytotoxicity of silyl- and carbonyl-substituted isoxazoles," <i>Chemistry of Heterocyclic Compounds</i> (2000) Vol. 36, No. 10, pp. 1226-1231	
	30	PLATE et al., "Synthesis and Muscarinic Activities of 3-(Pyrazolyl)-1,2,5,6-tetrahydropyridine Derivatives," <i>Bioorganic & Medicinal Chemistry Letters</i> (1996) Vol. 4, No. 2, pp. 227-237	
	31	Supplementary European Search Report dated February 6, 2009 for corresponding European Application No. 04788159.4	
	32	TANAKA et al., "An Effective Lewis Acid-Mediated 1,3-Dipolar Cycloaddition of Nitrile Oxide Using Acetylene: Synthesis of a (2-Aminopyridin-3-yl) isoxazole Derivative and its Application to Novel Antifungal Agents," pp. 1-8	
	33	VRZHESCHCH et al., "Supercooperativity in platelet aggregation: Substituted pyridyl isoxazoles, a new class of supercooperative platelet aggregation inhibitors," <i>FEBS Letters</i> (1994) Vol. 351, No. 2, pp. 168-170	

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